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AMERICAN CYANAMID CO

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New pyrimidine derivatives useful as herbicides, especially for selective weed control

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NOVELTY

Z-Aryloxy- or 2-arylitio-6-aryl pyrimidine derivatives (I) are new.

DETAILED DESCRIPTION

2-Aryloxy- or 2-arylphio-6-aryl-pyrimidine derivatives of formula are new.

C(7-D12, 14-V2) .2

A = optionally substituted anyl, optionally substituted 5- or 6membered heteroary) or diffuorobanzodioxoly);

B = phenyl or thisnyl;

m # 0-5:

 \mathbb{R}^3 = halomen. CN or optionally substituted alive, alkenyl, alkynyl. alkovyalkyi, haloalkyi, alkovy, halosikovy, alkyithio, alkylameno or dialkylämino:

 $\mathbb{R}^2 = \mathbb{H}$, halogan, CN or optionally substituted alkyl, altery, haloalkyl

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or haloalkovy;

 $R^3 = halogen$, NO_{20} CN, halosikyi, halosikony, halosikyithio, SF₂₀ or oprionally substituted alkyl, alkenyl, alkynyl, alkoxy, alkoxyalkyi, alkoxyalkoxy, alkyithio, alkyisuifinyi or aikvisuifonyi:

 $X \approx 0 \text{ or } S$.

Herbicidal. In a pre-emergence test, 2-(2-chloro-4-pyridyloxy)-6methyl-4-(4-mifluoromethylpitenyl)-pyrimidine at an application rate of 0.4 kg/ha gave 100% control of poppy (Paparer rhoeas) and 91-99% control of chickweed (Stellaria media).

MECHANISM OF ACTION

None given.

UME

(I) are herbicides useful for selective weed control, e.g. for preemergence weed control in winser wheat, maize, soya, conon or rice, or post-emergance weed control in winter wheat or maine.

ADVANTAGE

(I) have good selectivity and biodegradability. In a pre-emergence. test, 2-(2-chioro-4-pyridyioxy)-6-methyl-4-(4-trifinoromethylphenyl)pyrimidine at an application rate of 0.4 kg/ha caused no damage to winter wheat, maize, coya, cotton or rice.

SPECIFIC COMPOUNDS

9 Compounds (I) are specifically claimed, e.g. 4-methoxy-2-(1methyl-3-triflucromethyl-5-pyrazolyloxy)-6-(4millionomethylphenyl)-pyrimidine of formula (la).

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EXAMPLE

A mixture of 4-methyl-2-methylsalfonyl-6-(4trifluoromethylphenyl)-pyramidine (0.32 g), 3-trifluoromethylphenol (0.18 g), potassium carbonate (0.25 g) and acetonitrile (25 ml) was refluxed for 4 hours, diluted with water and extracted with methyl. acetate to give 4-mediyi-2-(3-trifluoromethyiphenyi)-6-(4diffisoromethy/phony!)-pyrimidine (0.39 g), m.pt. 124-127°C.

TECHNOLOGY FOCUS

Organic Chemistry - Preparation: (f) is prepared by:

(1) rescuing a compound of formula (III) with a metal compound of formula (IV) and oxidizing the product when L is hydrogen; or

(2) reacting a compound of formula (V) with a compound of formula

L = H or a leaving group; $M \approx Ci$, Mg, Za, B or Sa: Y = a leaving group; and M' = H or metal. (15pp367DwgNo.0/0)

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